

1 **Implicit adaptation is modulated by the relevance of feedback**

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12

13 **ABSTRACT**

14 Given that informative and relevant feedback in the real world is often intertwined with distracting and
15 irrelevant feedback, we asked how the relevancy of visual feedback impacts implicit sensorimotor
16 adaptation. To tackle this question, we presented multiple cursors as visual feedback in a center-out
17 reaching task and varied the task relevance of these cursors. In other words, participants were instructed
18 to hit a target with a specific task-relevant cursor, while ignoring the other cursors. In Experiment 1, we
19 found that reach aftereffects were attenuated by the mere presence of distracting cursors, compared to
20 reach aftereffects in response to a single task-relevant cursor. The degree of attenuation did not depend on
21 the position of the distracting cursors. In Experiment 2, we examined the interaction between task
22 relevance and attention. Participants were asked to adapt to a task-relevant cursor/target pair, while
23 ignoring the task-irrelevant cursor/target pair. Critically, we jittered the location of the relevant and
24 irrelevant target in an uncorrelated manner, allowing us to index attention via how well participants
25 tracked the position of target. We found that participants who were better at tracking the task-relevant
26 target/cursor pair showed greater aftereffects, and interestingly, the same correlation applied to the task-
27 irrelevant target/cursor pair. Together, these results highlight a novel role of task relevancy on modulating
28 implicit adaptation, perhaps by giving greater attention to informative sources of feedback, increasing the
29 saliency of the sensory prediction error.

30

31 **ACKNOWLEDGEMENTS**

32 This work was supported by grants R35 NS116883, R01 NS105839, and R01 NS1058389 from the
33 National Institutes of Health (NIH).

34 INTRODUCTION

35 Sensory feedback is continuously used to help keep that sensorimotor system calibrated, enabling
36 the production of accurate movements despite ongoing changes to one's body and environment. This
37 adaptive recalibration process is thought to be driven in part or whole by sensory prediction error (SPE),
38 the difference between the predicted and actual sensory feedback (Burge, Ernst, & Banks, 2008;
39 Chaisanguanthum, Joshua, Medina, Bialek, & Lisberger, 2014; Kording, Tenenbaum, & Shadmehr, 2007;
40 Mazzoni & Krakauer, 2006; Tsay, Haith, Ivry, & Kim, 2022; Wolpert, Ghahramani, & Jordan, 1995). In
41 this paper, we examine what happens to sensorimotor adaptation when the sensory feedback is complex,
42 ambiguous, and redundant.

43 When performing goal-oriented movements in the real world, the visual field is cluttered,
44 possibly obscuring the feedback of the movement (Burge et al., 2008; Körding & Wolpert, 2004; Tsay,
45 Avraham, et al., 2021; Tsay, Kim, Parvin, Stover, & Ivry, 2021; Wei & Körding, 2010) or even the target
46 itself (Meegan & Tipper, 1998). Furthermore, the motion of other visual objects may be sources of
47 distraction or interference. How does the motor system determine which signal is relevant for evaluating
48 the consequences of an action, and how is this process influenced by the presence of competing visual
49 signals?

50 Kasuga et al (2013) studied these questions using a center-out reaching task in which the
51 participants could not see their moving arm, but had to rely on feedback provided by a moving cursor
52 (Kasuga, Hirashima, & Nozaki, 2013). On most trials, a single cursor reflecting the participants true hand
53 position was presented. Interleaved with these were perturbation trials. On some of the perturbation trials,
54 the cursor was rotated relative to the true hand position by a variable angle, introducing a sensory
55 prediction error. On other perturbation trials, there were two or three cursors, each following a different
56 displaced trajectory. For example, one cursor might correspond to the true hand position with the other
57 cursor(s) rotated by varying amounts; or all of the cursors might be rotated from the true hand position.
58 Given that the number of cursors and their respective rotations was randomized across trials, implicit
59 adaptation was measured by calculating the trial-by-trial change in hand angle. The results showed that
60 the size of the trial-by-trial change could be modeled by taking the average predicted response to each of
61 the individual cursors, albeit with a global attenuation in comparison to the single cursor condition. Thus,
62 the response to two cursors, one at 0° (veridical feedback) and one rotated by 45° was similar to when
63 there were two rotated cursors, one at 15° and one at 30° . The fact that there was adaptation even in the
64 former case is especially surprising given that one cursor provided veridical feedback. The adaptation
65 system did not appear to be preferentially sensitive to veridical feedback.

66 One issue of note in the Kasuga study is that, from the participants' perspective, it may be
67 appropriate to produce a composite error signal from the individual cursors because all of the cursors
68 were potentially of equal importance and relevance to the task. However, in a natural environment, there
69 is likely to be one relevant source of feedback amongst irrelevant and potentially distracting sources of
70 information. To better understand how feedback drives sensorimotor adaptation, it is important to know
71 whether it is sensitive to the relevance of available feedback. In the context of online feedback control,
72 Reichenbach et al. (2014) provided a compelling demonstration that the motor system was indeed
73 sensitive to the relevancy of visual signals (Reichenbach, Franklin, Zatska-Haas, & Diedrichsen, 2014).
74 Participants performed reaching movements in which the feedback display included one cursor linked to
75 the true hand position as well as up to 4 distractor cursors that moved with a similar but predetermined,
76 and thus non-contingent velocity profile. At some point during the movement, one of the cursors made an
77 abrupt lateral shift. Rapid, online corrections to the perturbation were much stronger when the perturbed
78 cursor was the one linked to the hand position, compared to when the perturbed cursor was one of the
79 distractors. Similar effects of task relevance have been observed in force-field adaptation studies (Heald,
80 Ingram, Flanagan, & Wolpert, 2018), suggesting that implicit processes required for both on-line
81 corrections and sensorimotor adaptation are sensitive to the task-relevance of different feedback signals.

82 In contrast, other lines of research have highlighted how sensorimotor adaptation is seemingly
83 impervious to feedback regarding task goals and outcomes (Held & Gottlieb, 1958; Welch, 1969).
84 Consider the aiming landmark task, first introduced by Mazzone and Krakauer, (2006). After being briefly
85 exposed to a 45° visuomotor rotation, the participants were instructed to aim to a landmark positioned 45°
86 in the opposite direction from the target. By implementing this strategy, the participants were essentially
87 perfect after one trial, producing movements in which the rotated cursor hit the target. Nonetheless, over
88 the next 100 trials or so, performance deteriorated, with the hand angle increasing even further away from
89 the target (Taylor & Ivry, 2011). This paradoxical result arises because the adaptation system, impervious
90 to the strategy, recalibrates the sensorimotor mapping to reduce the SPE, here defined as the difference
91 between where the movement was directed (towards the landmark) and where the cursor appeared (at the
92 target). Similarly, participants implicitly adapt to the movement of a cursor that follows an invariant
93 spatial trajectory displaced from the target, even when they are fully aware of the manipulation and told to
94 ignore it (Avraham, Morehead, Kim, & Ivry, 2021; Kim, Morehead, Parvin, Moazzezi, & Ivry, 2018;
95 Morehead, Taylor, Parvin, & Ivry, 2017; Parvin, McDougle, Taylor, & Ivry, 2018; Tsay, Parvin, & Ivry,
96 2020).

97 Experiments such as these have led to the view that implicit adaptation is solely dependent on
98 SPE, impervious to feedback concerning the task outcome (Kim, Parvin, & Ivry, 2019). The results from
99 the multiple cursor study of Kasuga *et al.* (2013) would also be consistent with this hypothesis. However,
100 as noted above, the cursors were, in a sense, all task-relevant. Here we employ a multiple cursor task
101 similar to Kasuga *et al.*, but with two key changes. First, rather than randomize the perturbation from
102 trial-to-trial, we employed a fixed rotation throughout the training period and assessed adaptation in a
103 subsequent block where feedback was withheld. Second, and most importantly, we varied the cursor
104 configurations and instructions as a way to manipulate the task relevance of the different visual feedback
105 signals. In this way, we sought to determine whether implicit adaptation is sensitive to the relevance of
106 the feedback.

107

108 **METHODS**

109 **PARTICIPANTS**

110 Undergraduate students ($n = 64$, 41 females, age = 20 ± 2 years) were recruited from the
111 University of California, Berkeley community and financially compensated for their participation in the
112 experiment. All participants were right handed, as assessed by the Edinburgh Handedness Inventory
113 (Oldfield, 1971). The research protocol was approved by the UC Berkeley institutional review board.

114 **EXPERIMENTAL APPARATUS**

115 The participant was seated in front of a horizontally oriented computer monitor that was
116 supported by a table frame. All hand movements were tracked on a digitizing tablet (53.2 cm x 30 cm,
117 ASUS), positioned 27 cm below the monitor. The participant held a modified air hockey ‘paddle’
118 embedded with a digitizing stylus to make center-out reaching movements over the tablet surface in
119 response to visual stimuli displayed on the monitor. The participant’s hand was occluded by the
120 table/monitor, and the room was minimally lit to further preclude visual feedback of the arm. The latency
121 between the movement of the digitizing stylus and the updating of the cursor position on the monitor was
122 33 ms. The experimental code, controlling the visual display and acquisition of kinematic information
123 was written in MATLAB (version 2016), using the Psychophysics toolbox extensions (Pelli, 1997).

124 **OVERVIEW OF THE REACHING TASK**

125 Participants performed 8 cm reaches to targets located around a central starting location. The start
126 location was indicated by a 6 mm white annulus, and the target was a 6 mm blue circle. The visual

127 displays also included feedback cursors (3.5 mm white circles) that, depending on the condition, either
128 corresponded to the participant's hand position or were rotated around the start location at a fixed angle
129 from the hand position.

130 In all conditions, participants were instructed to produce rapid movements such that the task-
131 relevant designated cursor would 'shoot' through the target. Movement onset during the task was
132 arbitrarily defined as the time at which movement amplitude reached 1 cm from the center of the start
133 position. Movement time (MT) was defined as the duration from this point until the hand reached a radial
134 distance of 8 cm, the target distance. Auditory feedback concerning MT was used to encourage
135 participants to make relatively fast movements. For reaches shorter than 100 ms or longer than 300 ms,
136 the messages 'too fast' or 'too slow' were played over the computer speaker. A neutral 'knock' sound was
137 played if MT fell within the desired range. Across all trials, the median reaction time (RT) and MT were
138 462 ms and 172 ms, respectively. The median total trial time (TTT), defined as the time from the start of
139 one trial to the start of the next, was 3714 ms. One way ANOVAs revealed no differences between groups
140 for RT ($F_{(3, 44)} = 0.448, p = 0.720$), MT ($F_{(3, 44)} = 1.336, p = 0.275$), or TTT ($F_{(3, 44)} = 0.245, p = 0.865$).

141 On trials with visual feedback, the cursor or cursors were visible until the movement amplitude
142 reached 8 cm, whereupon the end point position was frozen for an additional 1 s. By freezing the cursors,
143 the participant received additional endpoint feedback of performance accuracy. At the end of the feedback
144 period, the cursors were turned off and the participant moved their hand back to the start position. To help
145 the participant find the start position, veridical feedback was provided when their hand was within 1 cm
146 of the start position. Once in the start position, the feedback cursor was turned off and the annulus filled,
147 indicating that the participant should prepare for the next trial. The next target appeared once the
148 participant remained within the starting position for 500 ms.

149 EXPERIMENT 1

150 Experiments 1a and 1b ($n = 48, 12$ per group) employed a similar design in which the participant
151 completed a series of five blocks. The *No Feedback Baseline* block was composed of 24 reaches without
152 visual feedback, one to each of 24 targets evenly spaced at 15° intervals (0° to 345°, with 0° corresponding
153 to a rightward movement). This block was included to familiarize the participants with the experimental
154 apparatus and with making movements in the desired time. The next block, *Feedback Baseline*, was
155 composed of 10 cycles of reaches, with each cycle composed of one reach to each of the 24 target
156 locations (240 trials). Veridical online feedback was provided by a feedback cursor aligned with the
157 participants' hand. Next was the *Training* block in which the specific experimental manipulations of the
158 cursor feedback were introduced (detailed below). Targets were limited to three locations (30°, 150°,
159 270°) with 80 cycles of 3-target sets (Training, 240 total trials). The 120° spacing was chosen to minimize
160 generalization/interference of adaptation effects between the three training locations (Day, Roemmich,
161 Taylor, & Bastian, 2016; Krakauer, Pine, Ghilardi, & Ghez, 2000). The *Aftereffect* block had one cycle of
162 24 trials with participants reaching to each target without visual feedback, similar to *No Feedback*
163 *Baseline*. At the start of this block, the participant was explicitly instructed to move their unseen hand
164 directly to the target. The *Washout* block had the same structure but with veridical visual feedback (3
165 cycles, or 72 trials). The *Aftereffect* and provided the critical data to test for aftereffects, indicative of
166 implicit adaptation and generalization. In all blocks, the order of the target location was randomized
167 within a cycle.

168 EXPERIMENT 1a

169 Experiment 1a was designed to examine the influence of distractor cursors on performance during
170 a visuomotor rotation task. We compared performance between training conditions in which the display
171 contained a single feedback cursor rotated 45° from the true hand position or when the display also
172 contained two additional cursors, positioned +/- 45° relative to the single cursor (Fig 1D).

173 *One Cursor group (n=12)*: A single feedback cursor was visible during the blocks with visual
174 feedback. In the *Feedback Baseline* and *Washout* blocks, the cursor provided veridical feedback. At the
175 start of the Training block, the participant was informed that the feedback cursor would no longer be
176 veridical but would now be displaced by 45° relative to their hand position (counterbalanced clockwise or
177 counterclockwise across participants). They were instructed that the task goal was to compensate for this
178 rotation such that the rotated cursor would hit the target. The experimenter explained the effect of the
179 rotation on the cursor and the new task by illustrating it on a whiteboard. Although it could be inferred
180 from the instructions that they could re-aim 45° away from the target to achieve the goal, we did not
181 explicitly instruct them to use such a strategy. By making the rotation explicit, we believed there would
182 be less ambiguity in the *Aftereffect* and *Washout* blocks, in which we assess implicit adaptation by
183 instructing participants to reach directly with their hand. Furthermore, since participants were informed
184 about the rotation of the cursors in the *Three Cursor group* (below), this made the number of cursors the
185 only difference between the two groups.

186 *Three Cursor group (n=12)*: Three feedback cursors were visible during each of the blocks with
187 visual feedback, with the cursors separated by 45°. Each cursor had a unique color: green, orange, and
188 purple (RGB values [255] [7 210 0], [231 145 53], [234 0 238] respectively). All colors were
189 approximately matched on luminance based on a ‘Hue Chrome Luminance’ color scheme, and the
190 assignment of color to cursor was counterbalanced across participants. In the *Feedback Baseline* and
191 *Washout* blocks, the participant was informed that the middle cursor would correspond to the true hand
192 position and instructed to hit the target with that cursor, specified in terms of the cursor color,
193 idiosyncratic for each participant. This cursor was flanked by the two distractor cursors, resulting in three
194 cursors moving at -45°, 0°, 45°, relative to hand position (Figure 1). The participant was instructed to
195 ignore the other two cursors. Thus, if the color assignment for the -45°, 0°, 45° cursors was green, orange,
196 and purple respectively, then the participant was to hit the target with the orange cursor and ignore the
197 green and purple cursors.

198 During the Training block, all three cursors were rotated by 45°, such that the cursors now
199 appeared at 0°, 45°, 90° (or 0°, -45°, -90°, counterbalanced across participants) relative to the true hand
200 position. The participant was informed of the manipulation and instructed to hit the target with the middle
201 cursor. Since the relative color assignment of the cursors remained the same, the instructions did not
202 change. Using the color mapping example from above with a counterclockwise rotation, the participant’s
203 goal was still to hit the target with the orange cursor (now rotated 45° from true hand position), while
204 ignoring the green and purple cursors. With this arrangement, the purple cursor now ended up
205 corresponding to the true hand position (0°) and the green cursor was rotated by 90°, relative to the hand.
206 As such, if each cursor contributes equally to form a composite SPE (Kasuga et al., 2013), the net SPE in
207 the *Three Cursor* condition is identical to that in the *One Cursor* condition.

208 EXPERIMENT 1b

209 As reported below, the inclusion of the two task-irrelevant distractors attenuated adaptation in the
210 *Three Cursor group*, relative to the *One Cursor group*. Experiment 1b was designed to test two hypotheses
211 that could account for this attenuation. The first hypothesis, “Error Averaging”, posits that adaptation is
212 equally driven by SPE signals generated from all three cursors, but that their weightings add up to less
213 than 1. Thus, the total amount of adaptation resulting from a 0°, 45°, and 90° would be less than from just
214 one 45° cursor. The second, “Non-Specific”, hypothesis is that the presence of distractors dilutes the
215 effects of adaptation in a general manner, and as such, the attenuation effect is not dependent on the
216 particular path of the distractor cursors. To evaluate these hypotheses, we compared two, three-cursor
217 variants in Experiment 1b, using the same trial structure as in Experiment 1a.

218 In the *Compensate group (n = 12)*, the visual feedback was not rotated during the Training block.
219 Instead, the participants were instructed to hit the target with the 45° side cursor. For example, if the
220 participant was in the clockwise group (again, counterbalanced across participants) and had the color

221 mapping shown in Figure 1D, they were instructed to hit the target with the green cursor. “Compensate”
222 refers to the fact that, while the three cursors were not rotated relative to baseline, the instructions
223 required participants to compensate for the angular offset of the task-relevant cursor. Note that as a result
224 of this change in instructions, the angular offset of task-relevant cursor from the hand position (by 45°) is
225 similar to the Three Cursor condition in Experiment 1a.

226 In the Ignore Rotation group ($n = 12$), the three cursors were rotated by 45° in the Training block.
227 However, unlike Experiment 1a, the task goal was changed for this block, with the participant instructed
228 to hit the target with the outer cursor that was in the opposite direction of the rotation (e.g., for a
229 counterclockwise rotation, the task-relevant cursor now became the cursor that was clockwise to the
230 center cursor). As shown in Fig. 1D, the net effect of the rotation and change in instructions results in the
231 task-relevant cursor corresponding to the position of the hand. “Ignore” here refers to the fact that
232 participants can ignore the 45° rotation applied to the three cursors since the task-relevant cursor ends up
233 being veridical with respect to the participant’s hand.

234 In summary, the groups used in Experiment 1a and 1b match average and task-relevant sensory
235 prediction errors across their conditions, offering a window into the unique contribution of these errors.
236 Furthermore, with the exception of the Ignore condition, all groups were required to make a 45° change in
237 reaching angle in the Training block to compensate for the perturbation, making reaching behavior very
238 similar across these groups.

239

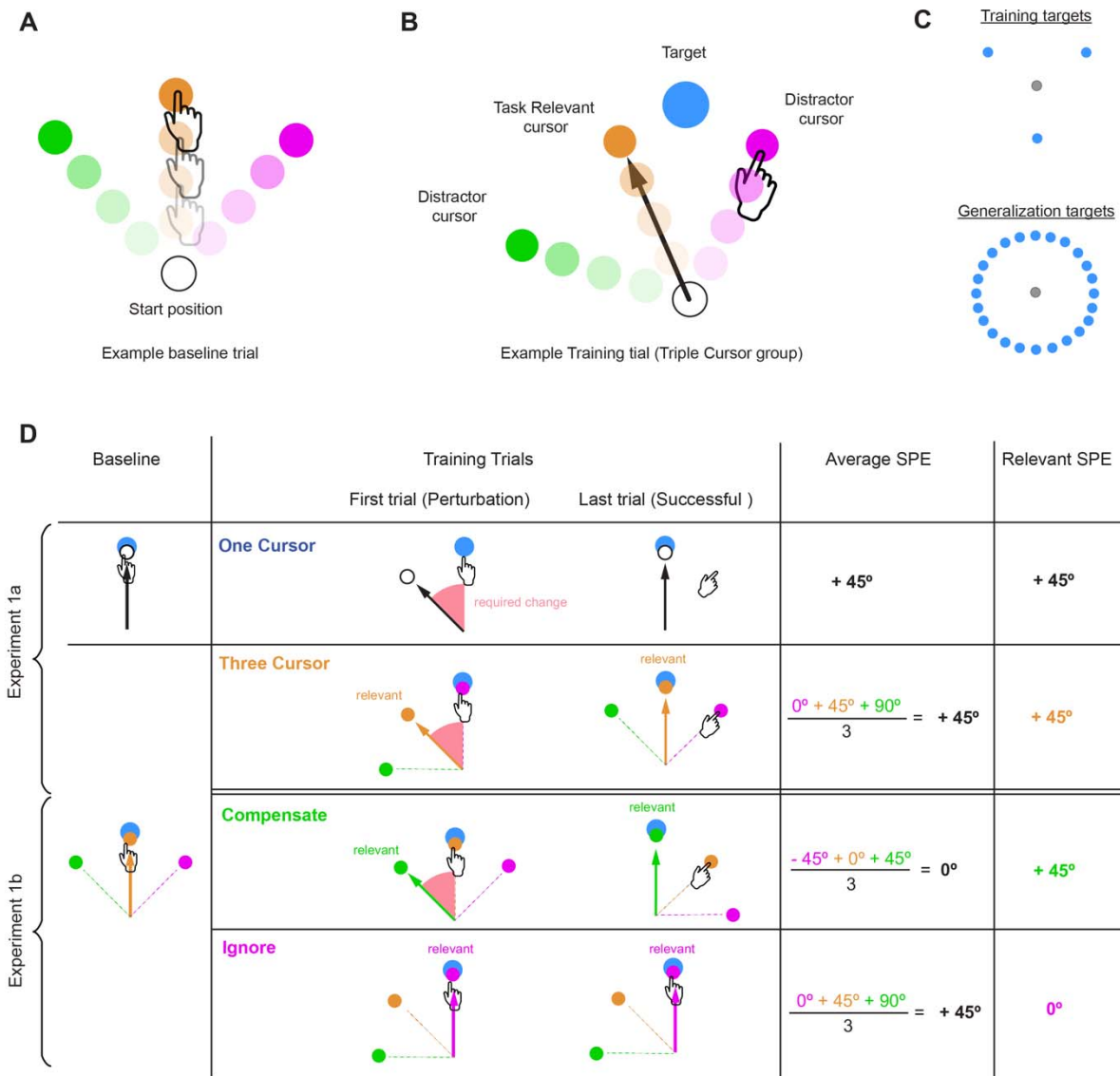
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246 **Figure 1: Feedback configurations for the Three Cursor group in Experiment 1.** **A)** In the baseline block, the middle cursor
 247 follows the veridical hand position, with two additional cursors rotated either +/- 45° on either side. Participants are told to reach
 248 directly to the target with the middle cursor. **B)** In the *Training* block, all the cursors are rotated by 45°, such that the middle
 249 cursor no longer corresponds to the true hand position. Participants are told to hit the target with the middle cursor and ignore the
 250 other two cursors. **C)** Target locations for Experiment 1. In the *Training* block, targets appeared in three locations (30°, 150°, 270°).
 251 In all other blocks, targets appeared in 24 generalized locations (from 0° to 345° in 15° intervals). **D)** Experimental
 252 conditions for Experiment 1. Experiment 1a compares learning from one cursor versus three cursors, keeping average SPE and
 253 relevant SPE the same. Experiment 1b employs two additional multiple cursor conditions to understand how the distractor
 254 cursors affect implicit adaptation. In the *Compensate* group, the three cursors were not rotated during the *Training* block, but the
 255 relevant cursor changed. Participants were instructed to hit with an outer cursor (green cursor), thus making that cursor relevant.
 256 This change in relevancy decreases the average SPE (i.e., 0°) but keeps the relevant SPE the same (i.e., 45°) as the *Three Cursor*
 257 group. In the *Ignore* group, the cursors were rotated by 45° during the *Training* block. Participants were told to hit the target with
 258 the cursor that followed their veridical hand position (pink cursor). Thus, the relevant cursor has change but the average SPE
 259 remains the same as the *Three Cursor* group.

260

261 EXPERIMENT 2

262 Experiment 2 ($n = 16$) was designed to further investigate how adaptation is influenced by task-
263 relevant feedback in the context of redundantly controlled objects. Generalization in this experiment was
264 assessed in one half of the workspace, with reaches limited to 13 target locations spaced every 15° (Fig.
265 4C). To control for workspace-specific training artifacts (e.g., biomechanical biases), participants were
266 assigned to reach in one of four areas of the workspace – Top (0° - 180°), Left (90° - 270°), Bottom (180° -
267 360°), or Right (270° - 90°). In the *No Feedback Baseline* block, the participant made three reaches to each
268 of the 13 targets without visual feedback. This was followed by the *Feedback Baseline* block, consisting
269 of five reaches to the 13 targets with veridical feedback provided by a single cursor.

270 In the *Training* block (200 trials, with a break at the midpoint), two targets were presented,
271 positioned approximately $\pm 45^\circ$ clockwise (see below) from the center of the participant's workspace.
272 Two cursors appeared during the reaching movement, one rotated clockwise (-45°) and the other
273 counterclockwise (45°) from the true hand position. The participant was instructed that one target-cursor
274 pair was task-relevant. For example, if the right target was task-relevant, then the participant's goal was to
275 hit this target with the cursor that was rotated in the clockwise direction. The participant was told to
276 ignore the other target-cursor pair. The side of the task-relevant target was counterbalanced across
277 participants. Note that by limiting the workspace to 180° , the instructions were stated in terms of a fixed
278 direction (e.g., "hit the target on the right" or "hit the upper target"). Following the *Training* block, the
279 participant completed an *Aftereffect* and *Washout* block (39 reaches each, three to each of 13 target
280 locations). These blocks provided the data for assessing aftereffects and generalization.

281 To verify that the participant was attempting to hit the task-relevant target, the location of each
282 target was independently jittered from trial to trial. The size of the jitter was one of five values, -10° , -5° ,
283 0° , 5° , and 10° , and determined in a pseudorandom manner for each participant, such that each value was
284 selected once every five trials. The overall order was pre-defined so that the correlation between the jitter
285 for the two targets would be zero across the experiment. As a result of ensuring they were uncorrelated,
286 the beta weights associated with each target were independent. By jittering the exact location of the target,
287 we could perform a multiple regression, using each target as a predictor of hand angle. The beta weights
288 obtained from this analysis indicate the participant's sensitivity to the relevant and distractor target.

289 DATA ANALYSIS

290 No statistical analyses were performed beforehand to determine sample size. The chosen sample
291 sizes were based on similar studies of visuomotor adaptation (Huang et al., 2011; Galea et al., 2015;
292 Vaswani et al., 2015), as well as considerations for counterbalancing. Experiment 1 ($n=12$ per group, 48
293 total) consisted of 2 participants for every cursor color combination (3) and direction (2, clockwise vs.
294 counterclockwise). Experiment 2 ($n= 16$) had 2 participants in every possible counterbalanced
295 configuration (4 workspaces x 2 directions).

296 The primary dependent variable of interest for all experiments was the heading angle of the hand.
297 This was defined as the signed angular difference between the position of the hand and the target angle at
298 peak radial velocity. Trials in which the hand angle was more than 90° away from the target were
299 excluded from the analysis (42 total, 0.15% of all trials over all participants). Excluding these trials did
300 not affect the outcome of any statistical tests. The data were averaged across cycles (24 successive
301 reaches; 1 reach to each target), and baseline subtracted to aid visualization. Baseline was defined as
302 mean hand angle over the last 3 movement cycles of the baseline phase with veridical feedback.

303 The degree of adaptation was quantified as the change in heading angle in the opposite direction
304 of the rotation. We calculated heading angle during early adaptation, late adaptation, and the aftereffect
305 phase. Following previous studies (Kim et al., 2018; Tsay, Avraham, et al., 2021), early learning was

306 defined as the mean hand angle over the cycles 3 – 7 to estimate the per trial rate of change during the
307 *Training* block. (We also performed a secondary analysis using cycles 1–10 and obtained nearly identical
308 results.). Late learning was defined as the mean hand angle over the last 10 cycles during the *Training*
309 block. The aftereffect was operationalized as the mean hand angle over all cycles of the *Aftereffect* block.

310 All analyses were conducted using custom scripts in Matlab (version 2016). Our experimental
311 design for Experiments 1a and 1b aimed to determine if there were group differences across adaptation
312 blocks (early adaptation, late adaptation, and aftereffect blocks). Accordingly, we performed two-tailed t-
313 tests to compare groups across these phases. We did not apply post-hoc corrections as our statistical tests
314 were based on a priori hypotheses regarding expected group differences; nonetheless, the key results
315 remain robust after applying Bonferroni corrections for the three comparisons. In Experiment 2, our
316 design focused on the effect of target relevance. We quantified this by correlating heading angles with
317 target angles; higher beta weights indicate greater attention to tracking the relevant/irrelevant jittered
318 cursor/target location. The beta weights were evaluated with one-sample t-tests to determine if they
319 differed from zero, and within-participant t-tests were used to compare beta weights across relevant and
320 irrelevant targets. We report standard measures of effect size (Cohen's d for between-participant
321 comparisons; Cohen's d_z for within-participant comparisons).

322 For the Gaussian fitting procedure in Experiments 1a and 1b, we used the 'fmincon' function in
323 MATLAB. We first fit the group average generalization curves using a Gaussian function characterized
324 by three free parameters subject to the following lower and upper bound constraints: standard deviation
325 $[0^\circ, 100^\circ]$, height $[0^\circ, 100^\circ]$, and mean $[-100^\circ, 100^\circ]$. To obtain 95% confidence interval estimates for
326 these parameters, we employed a bootstrapping procedure 1000 times. For each iteration, we resampled N
327 participants, where N represents the total number of participants in the experiment, with replacement. We
328 then calculated the average generalization function from this bootstrapped sample and derived the three
329 free parameters by fitting the Gaussian function.

330 For the Ignore group in Experiment 1b, we applied a cluster analysis approach to identify if there
331 were any significant clusters during training where the hand deviated from 0 (Tsay et al., 2020). This step
332 consisted of three steps. First, a t-test was performed for each cycle, asking if the observed hand angle
333 diverged from zero. Second, clusters were defined as epochs in which the p value from the t-tests were
334 less than 0.05 for at least two consecutive cycles. Third, to identify the probability of obtaining a cluster
335 of consecutive cycles with significant p values, we performed a permutation test. In this, we created 1000
336 permutations of the data with the cycles shuffled. For each shuffled permutation, we performed the first
337 two steps described above to identify clusters and for those meeting this criterion, we calculated the sum
338 of the t-scores over the significant cycles. Doing this for each of the 1000 permutations resulted in a
339 distribution of t-scores. The proportion of random permutations which resulted in a t-score of equal or
340 greater to that obtained from the data could therefore be directly interpreted as the p value.

341 Applying the first two steps to the actual data, we identified only one cluster, and only of length
342 two (training cycle 10 mean = 2.32° [$1.29^\circ, 3.35^\circ$], $t(11) = 4.956$, $p = 4.315e-4$, $d = 1.431$, and cycle 11
343 mean = 1.58° [$0.55^\circ, 2.62^\circ$], $t(11) = 3.362$, $p = 0.006$, $d = 0.971$). The sum of t-scores for the two
344 consecutive cycles ($t = 8.318$) was compared to the distribution generated by the permuted distribution,
345 producing a p-value of 0.004. We therefore concluded that this cluster of cycles represented a significant
346 deviation from 0, rather than being due to chance.

347 DATA AND CODE AVAILABILITY STATEMENT

348 Raw data and analysis code can be openly accessed at
349 https://github.com/DariusParvin/Adaptation_Multiple_Cursor_Experiments.

350

351 RESULTS

352 EXPERIMENT 1a

353 In Experiment 1, we examined the impact of task-irrelevant cursors on implicit adaptation. We
354 first compared adaptation to displays consisting of either a single cursor (One Cursor group) or three
355 cursors, separated by 45° (Three Cursor group, Fig 2). In both groups, participants were informed that,
356 during the training block, the task-relevant cursor (single or middle) was rotated by 45° relative to their
357 hand position, and that their task was to hit the target with this cursor. In the Three Cursor group,
358 participants were also told that one of the distractor cursors would coincide with their veridical hand
359 position but that they should ignore it and focus on the task-relevant cursor.

360 Performance of the two groups during the Training block was compared to assess effects of the
361 distractor cursors on overall learning. Both groups appeared to compensate for the perturbation at a
362 similar rate and extent. There were no significant differences in early learning (cycles 3-7, One Cursor
363 mean hand angle = 18.00° [9.90°, 26.10°], Three Cursor mean = 26.67° [17.23°, 36.12°], [$t_{(22)} = -1.535$, p
364 = 0.139, $d = -0.626$]), nor in late learning (last 10 cycles, One Cursor mean = 43.44° [41.27°, 45.62°],
365 Three Cursor mean = 41.55° [40.14°, 42.96°], [$t_{(22)} = 1.608$, $p = 0.122$, $d = 0.656$]). Previous studies have
366 shown that implicit adaptation is typically limited to 10°-25° of learning (Bond & Taylor, 2015; Kim et
367 al., 2018; Morehead et al., 2017; Tsay, Lee, Ivry, & Avraham, 2021). Given that the change in hand angle
368 is much larger than this range and that the participants were explicitly informed of the manipulation, we
369 assume there is a strategic (Tsay et al., 2023), aiming contribution to performance here. As such, the
370 results indicate that the distractor cursors did not have an appreciable influence on the participants' ability
371 to adopt an aiming strategy to complement adaptation.

372 To assess implicit adaptation, we included a block of trials after training in which the cursor was
373 no longer presented (Aftereffect block), and participants were asked to forgo strategy use and aim directly
374 to the target (i.e., "Move your hand straight to the target as fast and accurately as you can. Do not aim
375 away from the target"). Both groups exhibited aftereffects at the trained target location. However, the
376 magnitude of adaptation differed for the two groups: The One Cursor group had a significantly greater
377 aftereffect than the Three Cursor group (One Cursor group = 18.10° [14.50°, 21.69°], Three Cursor group
378 = 9.72° [6.48°, 12.96°], [$t_{(22)} = 3.808$, $p = 9.622e-4$, $d = 1.555$]).

379 There are at least two reasons why the aftereffect would be larger in the One Cursor group. First,
380 the difference could reflect that adaptation is attenuated by the inclusion of task-irrelevant distractors.
381 Alternatively, there could be a group difference in the use of an aiming strategy. Assuming that the
382 generalization function for adaptation is centered on the aiming location (Day et al., 2016), differences in
383 strategy use could cause differences in measured adaptation at the training location on aftereffect trials.

384 To evaluate these two hypotheses, we assessed generalization during the aftereffect block by
385 measuring adaptation across a set of probe targets that spanned the workspace in 15° increments (Figure
386 1C). The attenuation hypothesis predicts that the functions would be aligned but with a lower peak for the
387 Three Cursor group; the aiming hypothesis predicts that the functions would be similar in amplitude but
388 misaligned. The results were consistent with the attenuation hypothesis. While the generalization
389 functions for both groups were shifted towards the presumed aiming direction of aiming, the peak of the
390 generalization function was similar for the two groups. Based on parameters obtained when fitting a
391 Gaussian curve to the group data, the peak of the generalization function was 7.7° [4.4, 10.9] and 9.0°
392 [5.1, 12.8], with considerable overlap of the confidence intervals for the One Cursor and Three Cursor
393 groups, respectively [mean, bootstrapped 95% CI]. In contrast, the heights of the peaks were different for
394 the two groups (One Cursor: 18.8° [16.7, 20.8]; Three Cursor: 11.0° [9.3, 12.2]). These findings suggest
395 that the presence of the additional cursors attenuated the magnitude of implicit adaptation.

396 Error Averaging vs Relevant error models of adaptation

397 We next asked how the inclusion of the distractor cursors in the Three Cursor group influenced
398 adaptation. Specifically, we formulated two possible models to account for the attenuation.

399 The Error Averaging model, inspired by Kasuga and colleagues (Kasuga et al., 2013), proposes
400 that the error from each cursor is processed simultaneously and contributes towards learning (Equation 1).
401 By this model, the attenuation effect for the Three Cursor group is reflected in a reduction in the learning
402 term, B_t , relative to what it would be in the One Cursor case. Kasuga and colleagues observed a reduction
403 of 0.57 for B_t in their three cursor conditions relative to the One Cursor condition. This value is consistent
404 with our observed attenuation of 0.55 for the Three Cursor group in Experiment 1a (Three Cursor
405 aftereffect divided by the One Cursor aftereffect).

406

$$x_{n+1} = A * x_n + B_t * \frac{e_{relevant} + e_{distractor\ 1} + e_{distractor\ 2}}{3} \quad \text{Equation 1}$$

407

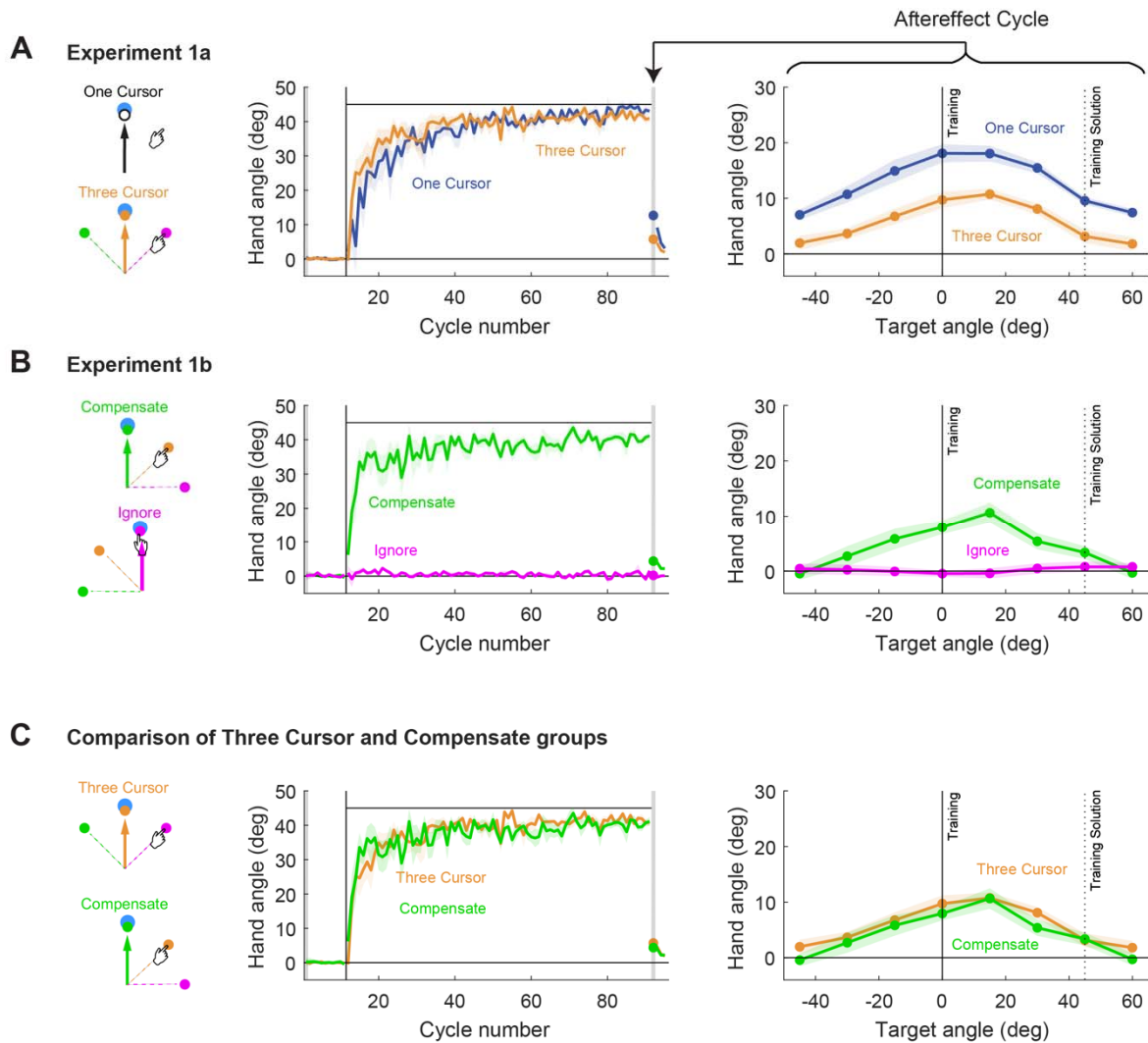
408 An alternative model, the Relevant Error model, proposes that implicit adaptation learns
409 selectively from the relevant error, while the distractor cursors attenuate adaptation in a non-specific
410 fashion. Thus, the directions of the distractor cursors have no bearing on the direction or magnitude of the
411 aftereffect. The attenuation due to the presence of the two distractor cursors would also be manifest in a
412 reduced B_t term, just as in the Error Averaging model (Equation 2).

413

$$x_{n+1} = A * x_n + B_t * e_{relevant} \quad \text{Equation 2}$$

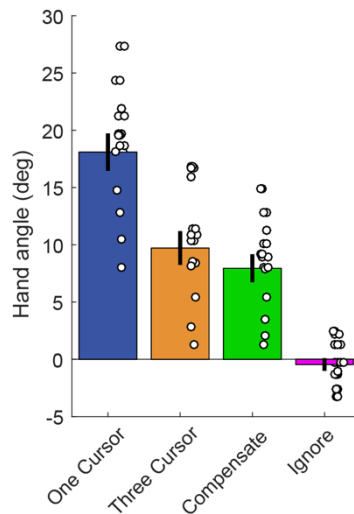
414

415



416

417 **Figure 2. Experiment 1 Results.** **A)** One Cursor and Three Cursor conditions in Experiment 1a. Middle: Both groups learn and
 418 compensate for the rotation at the same rate during training. Right: After the *Training* block, participants were instructed to reach
 419 directly to the generalization targets; no feedback was provided. The peak location of the generalization curves did not
 420 significantly differ, but the peak amplitude was greater in the One Cursor group compared to the Three Cursor group. **B)**
 421 Compensate and Ignore conditions in Experiment 1b. Middle: No change in hand angle was observed in the Ignore group. Right:
 422 Significant aftereffects were only seen in the Compensate group. **C)** Direct comparison of the Three Cursor and Compensate
 423 groups. No significant differences were observed in either the training (middle) or aftereffect (right) periods. Thick lines denote
 424 group means. Shaded regions denote \pm SEM.



425

426

427 **Figure 3.** Results from the aftereffect block in Experiment 1. Three Cursor and Compensate groups have similar levels of
428 aftereffects. Both groups also had significantly less than the One Cursor group, and significantly greater aftereffects than the
429 Ignore group. Error bars denote +/- SEM. Dots denote individual mean aftereffects.

430

431 EXPERIMENT 1b

432 To compare these two models, we tested two additional groups in Experiment 1b. The difference between
433 the two models is in their treatment of the distractor cursors. In the Error Averaging model, feedback
434 information from each cursor is equally weighted to form a composite error signal, whereas in the
435 Relevant Error model, only the feedback information from the task-relevant cursor is used to define the
436 error signal. Given that Error Averaging and Relevant Error models make identical predictions for the
437 Three Cursor group, we devised two, three-cursor variants that yield divergent predictions for the two
438 models.

439 In the Compensate group (Figure 2), the mapping between hand position and cursor is not
440 changed in the Training block; that is, the cursor that provided veridical feedback during the initial
441 Feedback block continues to provide veridical feedback in the Training block. However, the instructions
442 change, with the task goal now requiring the participant to hit the target with the cursor that is offset 45°
443 from the hand (45° clockwise or counterclockwise, counterbalanced across participants). To achieve this,
444 the participant must move in the opposite direction (e.g., clockwise if the task goal is to hit target with a
445 counterclockwise cursor). While the required hand trajectory is the same as in the Three Cursor group, the
446 set of errors is different. For the Compensate group, the sum is now 0° (-45°, 0°, 45°, relative to the hand
447 direction). Thus, the Error Averaging model would predict no adaptation. In contrast, the Relevant Error
448 model would predict the same amount of adaptation as observed with the Three Cursor group. Even
449 though the errors from the distractor cursors are different between the Compensate and Three Cursor
450 groups, both have a task-relevant cursor that is offset by 45° from the hand direction.

451 Consistent with the Relevant Error prediction, participants in the Compensate group exhibited a
452 robust aftereffect with Gaussian shaped generalization (mean at training location = 7.95° [5.26°, 10.63°];
453 one sample t-test against 0: $t_{(11)} = 6.510$, $p = 4.368e-5$, $d = 1.879$). Furthermore, this group behaved
454 similarly to the Three Cursor group during both the Training and Aftereffect blocks. There were no
455 significant differences during the Training block in their early learning (mean and 95% CI, Three Cursor

456 = 26.67° [17.22°, 36.11°], Compensate = 32.09° [25.70°, 38.48°], [$t_{(22)} = -1.047$, $p = 0.306$, $d = -0.428$],
457 nor in their late learning (mean and 95% CI, Three Cursor = 41.55° [40.14°, 42.96°], Compensate = 40.37°
458 [38.34°, 42.40°], [$t_{(22)} = 1.053$, $p = 0.304$, $d = 0.430$]). There were also no significant differences in their
459 aftereffects at the training locations (Three Cursor mean = 9.72° [6.48°, 12.96°], Compensate mean =
460 7.95° [5.26°, 10.63°], $t_{(22)} = 0.925$, $p = 0.365$).

461 The Ignore group provided a second test of the models. Here, a rotation of all three cursors was
462 introduced during the Training block, identical to that used for the Three Cursor group. However, the
463 participants were instructed in this block to hit the target with the 0° cursor, the one that was now
464 veridical with hand position. “Ignore” refers to the fact that the participant’s task was essentially to ignore
465 the rotation by focusing on the (new) cursor that matched their hand position. The Error Averaging model
466 predicts the same aftereffect as the Three Cursor group, since it is insensitive to the relevance of the error
467 signals and the average error is again 45° (90°, 45°, 0°, relative to the hand direction). The Relevant Error
468 model, on the other hand, predicts no aftereffect, since the hand direction and task-relevant cursor are
469 aligned. Again, the results of the aftereffects conformed to the Relevant Error prediction. A t-test showed
470 no significant aftereffect at the training location (mean = -0.45° [-1.66°, 0.75°]; one sample t-test against
471 0: $t_{(11)} = -0.833$, $p = 0.423$, $d = -0.240$).

472 The training period of the Ignore group lent itself to testing another prediction of the Error
473 Averaging hypothesis. If a composite SPE was present during the training trials, we should observe a
474 ‘drift’ away from the target, despite initial accurate performance (Mazzoni, 2006; Taylor and Ivry, 2011).
475 A t-test against 0 produced a non-significant result (mean = 0.57° [-0.20°, 1.33°], $t_{(11)} = 1.635$, $p = 0.130$, d
476 = 0.472). We were worried, however, that this measure would not be sufficiently sensitive to capture what
477 might be a transient effect (Taylor and Ivry, 2011). As a more sensitive alternative, we opted for a cluster
478 analysis, assessing if there were any consecutive cycles with significant drift. This approach identified a
479 significant cluster of two consecutive cycles (training cycles 10 and 11) in which the mean hand angle
480 was greater than 0° (mean = 1.95° [1.09°, 2.81°], $t_{(11)} = 4.992$, $p = 4.076e-4$, $d = 1.441$). While this small
481 cluster is in the expected direction, it is very small and of shorter duration than that observed in previous
482 studies (e.g., a 15° drift that lasts for about 80-100 trials, see Taylor & Ivry [2011]). Thus, if there was any
483 implicit adaptation in the ignore group, it was very limited.

484 Overall, the results rule out the Error Averaging model and are consistent with the Relevant Error
485 model: Significant aftereffects were observed when the task-relevant cursor was offset 45° from the hand
486 position during the Training block, either because we imposed a perturbation or altered the instructions.
487 Moreover, the results from the Ignore group, as well as the similarities between the Compensate and the
488 Three Cursor group, suggest there is minimal, or no specific influence from the distractor cursors. Rather,
489 the presence of the distractors appears to have a non-specific attenuation effect on adaptation to the task-
490 relevant cursor.

491

492 EXPERIMENT 2

493 Although the results of Experiment 1 suggest that task relevance modulates adaptation, they are
494 far from conclusive in this regard. Not only was there no imposed rotation for the Ignore group, just a
495 change in the task-relevant color in the Training block, this group also did not have to aim away from the
496 target (as in the Compensate group). To provide a more direct assessment of the contribution of task-
497 irrelevant information, we conducted a second experiment, using a design in which there were two targets
498 positioned approximately 90° apart, and two associated cursors, one rotated 45° in the clockwise direction
499 from the hand direction and the other rotated 45° in the counterclockwise direction (Figure 4A). One
500 target/cursor was designated task-relevant, and the other was designated a distractor (i.e., task-irrelevant).
501 Regardless of which was task-relevant, the movement required to hit the target was directed around the
502 midpoint of the two targets, resulting in each cursor landing near its associated target. We measured the

503 aftereffects around each target location to determine the degree of adaptation. We hypothesized that the
504 task-relevant target may form the locus of generalization for implicit adaptation. Therefore, whether there
505 is also an aftereffect around the distractor target was of primary interest.

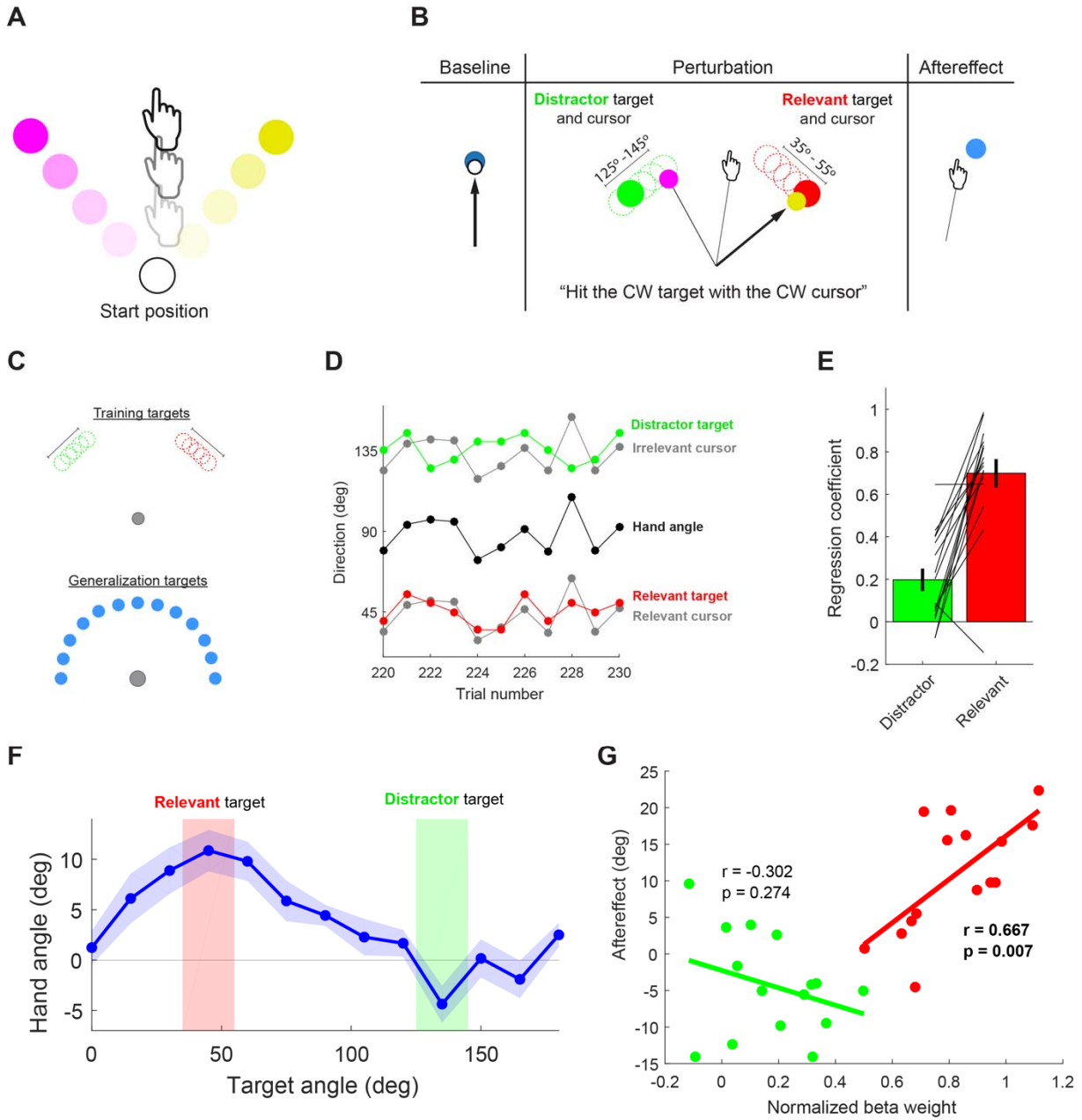
506 To verify that participants were following the instructions, we jittered the exact position of the
507 targets (see Methods) and calculated the trial-by-trial hand angle relative to each target (Figure 4D).
508 Using these time series, we then performed a multiple regression using the positions of the relevant and
509 distractor targets on each trial to predict the hand angle. This analysis produced a beta weight for each
510 target, quantifying how much the participant's behavior reflected the tracking of each target (Figure 4E).
511 We saw that greater weight was given to the task-relevant target in 15 of the 16 participants. The other
512 participant appeared to not follow directions (beta weights were -0.14 and 0.09 for the relevant and
513 distractor targets), and this person's data were excluded from the remaining analyses.

514 The mean beta weight for the task-relevant target was 0.76 [0.67, 0.84] and for the distractor
515 target was 0.20 [0.09, 0.32]. These values were significantly different from one another ($t_{(28)} = 8.149$, $p =$
516 $7.16e-9$, $d = 2.975$). Interestingly, the beta weight for the task distractor target was significantly greater
517 than zero ($t_{(14)} = 3.675$, $p = 0.002$, $d = 0.949$). Thus, while the participants followed the instruction to
518 track the relevant target, the results indicate that the participants were not able to completely ignore the
519 distractor target. One possibility is that the distractor target may have served as an additional visual
520 reference for aiming.

521 To assess adaptation, participants made reaching movements without feedback to an array of
522 targets, spanning the workspace that encompassed the two positions of the targets during the training
523 block. As in Experiment 1, the participants were instructed to reach directly towards each target. To
524 analyze these data, we collapsed across the two groups (clockwise or counterclockwise rotation of the
525 task-relevant cursor), displaying the data as though all participants were instructed to hit the clockwise
526 target and ignore the counterclockwise target (Figure 4B). A significant aftereffect in the direction
527 consistent with the rotation was observed around the task-relevant target location (mean 10.87° [6.48°,
528 15.27°]; one sample t-test against 0: $t_{(14)} = 5.304$, $p = 1.114e-4$, $d = 1.370$). Interestingly, there was also a
529 significant aftereffect at the distractor target location (mean -4.37° [-8.27°, -0.47°], $t_{(14)} = -2.406$, $p =$
530 0.031 , $d = -0.621$). The negative aftereffect associated with the distractor target was consistent with the
531 distractor cursor having the opposite rotation to the relevant cursor. When we directly compared the
532 aftereffects across the two targets by flipping the sign of distractor cursor, aftereffects were marginally
533 greater in the task-relevant target compared to the distractor cursor ($t_{(14)} = 2.00$, $p = 0.067$) with a medium
534 effect size ($d = 0.51$).

535 The fact that we observed a small aftereffect around the distractor target location would seem at
536 odds with the results of Experiment 1 where the effect of the distractor cursors was not dependent on their
537 direction (e.g., non-specific attenuation). However, as shown in the beta weight analysis, the participants
538 in Experiment 2 did not completely ignore the distractor target. To look at the relationship between
539 tracking and adaptation, we examined the correlation between beta weights and magnitude of the
540 aftereffect for both the relevant and distractor locations. To capture how much a participant weighted the
541 relevant target over the distractor, we normalized the beta weights (beta weight for relevant target / sum
542 of both beta weights) in these correlations. For the relevant location, the correlation was significant ($r =$
543 0.67 $p = 0.007$), meaning that those who were most successful in tracking the relevant target had the
544 largest aftereffect around that target's associated location. We performed the same analysis for the
545 distractor target. Since the distractor cursor had the opposite rotation, the predicted correlation would be
546 negative. This analysis revealed a similar pattern, but the correlation was not significant ($r = -0.30$, $p =$
547 0.274). We note that this analysis may be underpowered, given that the range both aftereffect and beta
548 weights are narrower for the distractor location compared to the relevant location.

549



550 **Figure 4. Results of Experiment 2.** **A)** Feedback configuration for training trials. Two cursors appeared during the reach, rotated
551 $\pm 45^\circ$ to the veridical hand position. **B)** Experimental task. Baseline and aftereffect trials had one-cursor feedback reaches to
552 generalization targets. During training, one target/cursor pair was made task-relevant by instructing the participant to focus on
553 hitting that target. By jittering the targets, we can assess how well the participant tracked the relevant target. **C)** Target locations,
554 limited to one half of the workspace. Top: Training target locations, centered $\pm 45^\circ$ from the midpoint of the workspace. The
555 exact location of each target was independently jittered by one of 5 values. Bottom, 13 generalization targets spaced 15° apart. **D)**
556 Trial by trial tracking behavior for distractor (green) and relevant (red) target/cursor pairs. **E)** Beta weights for the relevant target
557 (red) are significantly greater than for the distractor target (green). Data are group mean \pm SEM with lines representing
558 individual performance. **F)** Generalization functions. The magnitude of the aftereffect is larger around the relevant target location
559 (red) compared to the irrelevant distractor location (green), although the latter is also significantly greater than zero. **G)**
560 Correlation between beta weights and aftereffect for relevant and distractor targets. Significant correlation for the relevant target
561 (red) shows that participants who were better at tracking also had larger aftereffects. Non-significant correlation for distractor
562 target (green). Note that in Panels D and F, the data are transformed to graphically depict the relevant target at 45° for all
563 participants.

564 **DISCUSSION**

565 In this study, we asked how implicit adaptation, an important process for maintaining calibration
566 within the sensorimotor system, is affected by the presence of multiple visual signals. Through our use of
567 task instructions, we varied the information value of the signals, designating one cursor as task-relevant
568 and the others as task-irrelevant. As expected, we found that adaptation was sensitive to the rotation of the
569 task-relevant cursor. Moreover, as shown in Experiment 2, the degree to which participants tracked the
570 target with the relevant cursor predicted the size of their aftereffect, pointing to a strong relationship
571 between the task relevance assigned to the cursor and the amount of adaptation. Interestingly, implicit
572 adaptation was attenuated by the presence of distracting feedback, but this effect seemed to be non-
573 specific: The presence of the distractor cursors reduced the magnitude of adaptation, but not the direction
574 of adaptation. These findings highlight a novel role of task relevance for implicit adaptation.

575 By presenting multiple cursors and manipulating their relevance, we were able to make sense of
576 two seemingly contradictory findings in the sensorimotor adaptation literature. The results from several
577 studies suggest the motor system is sensitive to the relevance of feedback (Heald et al., 2018;
578 Reichenbach et al., 2014). In contrast, the results from other studies have shown that implicit adaptation is
579 insensitive to task goals. For example, when only one cursor is presented, implicit adaptation will learn
580 from the SPE, regardless of whether participants are explicitly told to ignore the visual feedback, or even
581 when the response to that cursor is detrimental to performance (Mazzoni & Krakauer, 2006; Morehead et
582 al., 2017; Taylor & Ivry, 2011; Taylor, Klemfuss, & Ivry, 2010). Our results suggest a hybrid position:
583 When more than one source of visual motion feedback is present, the primary input to the adaptation
584 system is the most relevant source of visual motion feedback. This selectivity constraint has been
585 observed in studies of other implicit motor functions, such as the finding that online corrections are faster
586 in response to perturbations of a task-relevant cursor (Reichenbach et al., 2014) or that separate force
587 fields can be learned depending on what part of a virtual tool is deemed relevant (Heald et al., 2018).

588 Although adaptation was sensitive to the relevance of the feedback, the overall attenuation of
589 learning demonstrated that the selectivity was imperfect. The presence of multiple cursors had a similar
590 attenuating effect in the present study as observed in Kasuga et al. (2013). Specifically, in both studies,
591 adaptation was attenuated by about 45% when there were three cursors relative to a standard single cursor
592 condition. This similarity is striking given that Kasuga used unpredictable rotations, interleaved the single
593 and multi-cursor conditions, provided no instructions concerning relevance, and measured learning using
594 a trial-by-trial, whereas we used a predictable rotation, blocked the conditions, instructed the participants
595 to attend one one cursor, and measured learning in an aftereffect block. An attenuation of motor responses
596 due to the presence of irrelevant feedback has also been observed for online corrections to cursor jumps
597 (Reichenbach et al., 2014).

598 The Relevance Model posits that the attenuation is non-specific; it could be that the irrelevant
599 information diverts visual attention away from the relevant cursor, thereby reducing adaptation.
600 Previously, Taylor and Thoroughman (2007) demonstrated that participants adapted less to perturbations
601 in a force field reaching adaptation task when they were engaged in a secondary task designed to divide
602 their attention (Taylor & Thoroughman, 2007). Given the similarity between the cursors, namely that all
603 originated from the same start position and moved with the same spatiotemporal correlation, it is likely
604 that the irrelevant cursors resulted in some degree of attentional diversion in our task (Folk, Remington, &
605 Johnston, 1992). This attentional explanation is also relevant when considering why participants are
606 sensitive to task-relevance in the presence of multiple sources of feedback yet are unable to ignore a
607 single cursor when instructed to do so.

608 However, the effect of visual distractors may be more nuanced than the non-specific attenuation
609 we have posited. Specifically, in Experiment 1b, the visual distractors in the Ignore condition elicited
610 negligible implicit adaptation, whereas in Experiment 2, the visual distractors elicited significant
611 aftereffects around the irrelevant target locations. We attribute this discrepancy to at least two factors:

612 First, the presence of a veridical cursor and its associated target may modulate the degree to which the
613 system adapts to irrelevant visual errors. In the Ignore condition of Experiment 1b, where both the
614 veridical cursor and an aiming target are present, participants may fully attend to the veridical
615 cursor/target pair, thus eliminating implicit adaptation in response to the irrelevant visual errors. In
616 Experiment 2, when neither a veridical cursor nor aiming target are provided, participants may lack a
617 clear referent for their actual hand position (Synofzik, Thier, Leube, Schlotterbeck, & Lindner, 2010;
618 Tsay, Kim, et al., 2021; Tsay, Kim, Haith, & Ivry, 2022). Here the system may not be able to fully
619 dismiss the irrelevant cursor/target pair, resulting in a modest degree of adaptation near the irrelevant
620 cursor/target location.

621 Second, the characteristics of the irrelevant visual errors may also modulate the degree to which
622 this information is ignored. In Experiment 1b, the irrelevant visual errors were large and introduced
623 abruptly. This may have led them to be easily deemed irrelevant, being outside the typical range of motor
624 noise (Wei & Körding, 2009), and thus, not sufficient to elicit implicit adaptation. In contrast, in
625 Experiment 2 the visual errors were small and were introduced more gradually, features that have made
626 them be deemed as more relevant by the adaptation system (Ingram et al., 2000; Kagerer, Contreras-
627 Vidal, & Stelmach, 1997). Future studies are required to systematically examine the characteristics that
628 influence how the nervous system evaluates error relevance, and in turn, how this perceived relevance
629 impacts the extent of implicit adaptation (e.g., (Makino, Hayashi, & Nozaki, 2023)).

630 **Limitations and future directions**

631 A limitation of our study is that we did not control or monitor fixation. Since it is likely that
632 participants directed their gaze at the relevant target (Neggers & Bekkering, 2000), one might argue that
633 the dominance of the relevant cursor in driving adaptation could reflect the fact that distractor information
634 is presented at more peripheral locations(s). We think that a fixation-based argument is unlikely to
635 account for the effects observed here. First, the distractor cursors attenuated adaptation in all conditions,
636 including in Experiment 2 where the relevant and distractor cursors were 90° apart. Thus, at least for the
637 angular separations employed here, there is no obvious relationship between distance and attenuation.
638 Second, prior work (Rand & Rentsch, 2015) has shown that magnitude of implicit adaptation is similar if
639 participants are required to maintain fixation on the start position, target, or allowed to gaze freely.

640 Although the present work makes clear that adaptation is impacted when operating in a complex
641 environment, one that is more akin to our natural environments, the processing stage at which this
642 interaction occurs is unclear. The present work suggests the operation of a selection process that
643 constrains the sensory prediction error, with the strength of this signal attenuated in the presence of task-
644 irrelevant information. Alternatively, the presence of uncertainty may operate as some sort of gain on
645 adaptation, either in terms of the strength of the sensory prediction or weight given to the sensory
646 prediction error. An important question for future research is to elucidate the processes involved in the
647 differential weighting of relevant and irrelevant sensory feedback, and specify how this information
648 impacts adaptation.

649 **References**

- 650 Avraham, G., Morehead, R., Kim, H. E., & Ivry, R. B. (2021). Reexposure to a sensorimotor perturbation
651 produces opposite effects on explicit and implicit learning processes. *PLoS Biology*, *19*(3),
652 e3001147.
- 653 Bond, K., & Taylor, J. A. (2015). Flexible explicit but rigid implicit learning in a visuomotor adaptation
654 task. *Journal of Neurophysiology*, *113*(10), 3836–3849.
- 655 Burge, J., Ernst, M. O., & Banks, M. S. (2008). The statistical determinants of adaptation rate in human
656 reaching. *Journal of Vision*, *8*(4), 20.1–19.
- 657 Chaisanguanthum, K. S., Joshua, M., Medina, J. F., Bialek, W., & Lisberger, S. G. (2014). The neural
658 code for motor control in the cerebellum and oculomotor brainstem. *ENeuro*, *1*(1),
659 ENEURO.0004-14.2014.
- 660 Day, K. A., Roemmich, R. T., Taylor, J. A., & Bastian, A. J. (2016). Visuomotor Learning Generalizes
661 Around the Intended Movement. *ENeuro*, *3*(2). doi:10.1523/ENeuro.0005-16.2016
- 662 Folk, C. L., Remington, R. W., & Johnston, J. C. (1992). Involuntary covert orienting is contingent on
663 attentional control settings. *Journal of Experimental Psychology. Human Perception and*
664 *Performance*, *18*(4), 1030–1044.
- 665 Heald, J. B., Ingram, J. N., Flanagan, J. R., & Wolpert, D. M. (2018). Multiple motor memories are
666 learned to control different points on a tool. *Nature Human Behaviour*, *2*(4), 300–311.
- 667 Held, R., & Gottlieb, N. (1958). Technique for studying adaptation to disarranged hand-eye coordination.
668 *Perceptual and Motor Skills*, *8*(3), 83–86.
- 669 Ingram, H. A., van Donkelaar, P., Cole, J., Vercher, J. L., Gauthier, G. M., & Miall, R. C. (2000). The
670 role of proprioception and attention in a visuomotor adaptation task. *Experimental Brain*
671 *Research. Experimentelle Hirnforschung. Experimentation Cerebrale*, *132*(1), 114–126.
- 672 Kagerer, F. A., Contreras-Vidal, J. L., & Stelmach, G. E. (1997). Adaptation to gradual as compared with
673 sudden visuo-motor distortions. *Experimental Brain Research. Experimentelle Hirnforschung.*
674 *Experimentation Cerebrale*, *115*(3), 557–561.
- 675 Kasuga, S., Hirashima, M., & Nozaki, D. (2013). Simultaneous processing of information on multiple
676 errors in visuomotor learning. *PloS One*, *8*(8), e72741.
- 677 Kim, H. E., Morehead, R., Parvin, D. E., Moazzezi, R., & Ivry, R. B. (2018). Invariant errors reveal
678 limitations in motor correction rather than constraints on error sensitivity. *Communications*
679 *Biology*, *1*, 19.
- 680 Kim, H. E., Parvin, D. E., & Ivry, R. B. (2019). The influence of task outcome on implicit motor learning.
681 *ELife*, *8*. doi:10.7554/eLife.39882
- 682 Kording, K. P., Tenenbaum, J. B., & Shadmehr, R. (2007). The dynamics of memory as a consequence of
683 optimal adaptation to a changing body. *Nature Neuroscience*, *10*(6), 779–786.
- 684 Körding, K. P., & Wolpert, D. M. (2004). Bayesian integration in sensorimotor learning. *Nature*,
685 *427*(6971), 244–247.
- 686 Krakauer, J., Pine, Z. M., Ghilardi, M. F., & Ghez, C. (2000). Learning of visuomotor transformations for
687 vectorial planning of reaching trajectories. *The Journal of Neuroscience: The Official Journal of*
688 *the Society for Neuroscience*, *20*(23), 8916–8924.
- 689 Makino, Y., Hayashi, T., & Nozaki, D. (2023). Divisively normalized neuronal processing of uncertain
690 visual feedback for visuomotor learning. *Communications Biology*, *6*(1), 1286.
- 691 Mazzoni, P., & Krakauer, J. W. (2006). An implicit plan overrides an explicit strategy during visuomotor
692 adaptation. *The Journal of Neuroscience*, *26*(14), 3642–3645.
- 693 Meegan, D. V., & Tipper, S. P. (1998). Reaching into cluttered visual environments: spatial and temporal
694 influences of distracting objects. *The Quarterly Journal of Experimental Psychology. A, Human*
695 *Experimental Psychology*, *51*(2), 225–249.
- 696 Morehead, R., Taylor, J. A., Parvin, D. E., & Ivry, R. B. (2017). Characteristics of Implicit Sensorimotor
697 Adaptation Revealed by Task-irrelevant Clamped Feedback. *Journal of Cognitive Neuroscience*,
698 *29*(6), 1061–1074.

- 699 Neggers, S. F., & Bekkering, H. (2000). Ocular gaze is anchored to the target of an ongoing pointing
700 movement. *Journal of Neurophysiology*, 83(2), 639–651.
- 701 Oldfield, R. C. (1971). The assessment and analysis of handedness: the Edinburgh inventory.
702 *Neuropsychologia*, 9(1), 97–113.
- 703 Parvin, D. E., McDougle, S. D., Taylor, J. A., & Ivry, R. B. (2018). Credit Assignment in a Motor
704 Decision Making Task Is Influenced by Agency and Not Sensory Prediction Errors. *The Journal*
705 *of Neuroscience: The Official Journal of the Society for Neuroscience*, 38(19), 4521–4530.
- 706 Pelli, D. G. (1997). The VideoToolbox software for visual psychophysics: transforming numbers into
707 movies. *Spatial Vision*, 10(4), 437–442.
- 708 Rand, M. K., & Rentsch, S. (2015). Gaze locations affect explicit process but not implicit process during
709 visuomotor adaptation. *Journal of Neurophysiology*, 113(1), 88–99.
- 710 Reichenbach, A., Franklin, D. W., Zatka-Haas, P., & Diedrichsen, J. (2014). A dedicated binding
711 mechanism for the visual control of movement. *Current Biology: CB*, 24(7), 780–785.
- 712 Synofzik, M., Thier, P., Leube, D. T., Schlotterbeck, P., & Lindner, A. (2010). Misattributions of agency
713 in schizophrenia are based on imprecise predictions about the sensory consequences of one's
714 actions. *Brain: A Journal of Neurology*, 133(Pt 1), 262–271.
- 715 Taylor, J. A., & Ivry, R. B. (2011). Flexible cognitive strategies during motor learning. *PLoS*
716 *Computational Biology*, 7(3), e1001096.
- 717 Taylor, J. A., Klemfuss, N. M., & Ivry, R. B. (2010). An explicit strategy prevails when the cerebellum
718 fails to compute movement errors. *Cerebellum*, 9(4), 580–586.
- 719 Taylor, J. A., & Thoroughman, K. A. (2007). Divided attention impairs human motor adaptation but not
720 feedback control. *Journal of Neurophysiology*, 98(1), 317–326.
- 721 Tsay, J. S., Avraham, G., Kim, H. E., Parvin, D. E., Wang, Z., & Ivry, R. B. (2021). The Effect of Visual
722 Uncertainty on Implicit Motor Adaptation. *Journal of Neurophysiology*.
723 doi:10.1152/jn.00493.2020
- 724 Tsay, J. S., Haith, A. M., Ivry, R. B., & Kim, H. E. (2022). Interactions between sensory prediction error
725 and task error during implicit motor learning. *PLoS Computational Biology*, 18(3), e1010005.
- 726 Tsay, J. S., Kim, H. E., McDougle, S. D., Taylor, J., Haith, A., Avraham, G., ... Ivry, R. (2023). *Strategic*
727 *Processes in Sensorimotor Learning: Reasoning, Refinement, and Retrieval*. Retrieved from
728 psyarxiv.com/x4652
- 729 Tsay, J. S., Kim, H. E., Parvin, D. E., Stover, A. R., & Ivry, R. B. (2021). Individual differences in
730 proprioception predict the extent of implicit sensorimotor adaptation. *Journal of*
731 *Neurophysiology*. doi:10.1152/jn.00585.2020
- 732 Tsay, J. S., Kim, H., Haith, A. M., & Ivry, R. B. (2022). Understanding implicit sensorimotor adaptation
733 as a process of proprioceptive re-alignment. *eLife*, 11. doi:10.7554/eLife.76639
- 734 Tsay, J. S., Lee, A. S., Ivry, R. B., & Avraham, G. (2021). Moving outside the lab: The viability of
735 conducting sensorimotor learning studies online (p. 2021.01.30.181370).
736 doi:10.1101/2021.01.30.181370
- 737 Tsay, J. S., Parvin, D. E., & Ivry, R. B. (2020). Continuous reports of sensed hand position during
738 sensorimotor adaptation. *Journal of Neurophysiology*, 124(4), 1122–1130.
- 739 Wei, K., & Körding, K. (2009). Relevance of error: what drives motor adaptation? *Journal of*
740 *Neurophysiology*, 101(2), 655–664.
- 741 Wei, K., & Körding, K. (2010). Uncertainty of feedback and state estimation determines the speed of
742 motor adaptation. *Frontiers in Computational Neuroscience*, 4, 11.
- 743 Welch, R. B. (1969). Adaptation to prism-displaced vision: The importance of target-pointing. *Perception*
744 *& Psychophysics*, 5(5), 305–309.
- 745 Wolpert, D. M., Ghahramani, Z., & Jordan, M. I. (1995). An internal model for sensorimotor integration.
746 *Science*, 269(5232), 1880–1882.